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UNITED STATES PATENT APPLICATION
OF
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FOR
PARTICLE SIZER AND DNA SEQUENCER

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PARTICLE SIZER AND DNA SEQUENCER

1. FIELD OF THE INVENTION

The present invention relates to an apparatus and method for distinguishing the size of a particle. More specifically, the invention relates to apparatuses and methods for screening the relative size of a particle within a capillary.

2. TECHNICAL BACKGROUND

Microbes are a common cause of illness and death. In order for medical professionals to diagnose and treat an illness it is often necessary to identify the disease causing agent. Moreover, identifying the cause of an outbreak of disease allows steps to be taken to prevent the spread of the disease.

Microbes are sometime identified based on physical characteristics such as size, shape, colony type, and staining characteristics. While such physical characteristics can be used to eliminate some of the possible disease causing agents, it is often not possible to identify a microbe based on these characteristics alone. For example many species of bacteria are further classified into strains. Some members of the species can be harmless or even beneficial microbes, while others may be pathogenic.

Because the disease causing strains have a gene or genes that render them pathogenic, DNA analysis can be used to identify these strains. The DNA analysis can provide a positive identity of a pathogenic organism. However, the amount of information provided is limited, and the process can take a considerable amount of time.

1 When the purpose of identifying the microbe is to treat a disease or prevent its spread the
2 DNA analysis may take too long to be useful.

3
4 Amplified Fragment Length Polymorphism (AFLP) and strain-specific
5 polymerase chain reaction (PCR) analyses are the methods used for determining the
6 identity of microbes. These procedures provide significantly more information than
7 standard DNA analysis and are more rapid and less expensive than extensive DNA
8 sequencing. AFLP analysis can be used for rapidly characterizing unknown pathogenic
9 species and strains, thereby providing valuable information for developing a therapeutic
10 response to an outbreak thereof. Strain-specific PCR analysis can rapidly identify a
11 previously studied threat species, even if the sample is present in a complex sample
12 mixture. Both methods require further analysis of the reaction results to determine the
13 size of the DNA fragments. Simple detection of the presence or absence of the fragment
14 is insufficient. The time required to complete the reaction analysis is presently between
15 15 and 20 minutes. However, size analysis is currently conducted using an automated
16 DNA sequencer. The gel procedure requires approximately three hours to complete, plus
17 additional time to set up the sequencing unit and download the resulting data. When
18 faced with the potential or actual release of a biothreat agent, it is important to obtain the
19 genetic information about the released organism in a significantly shorter time period. It
20 is also important to be able to conduct analyses in a field laboratory using affordable
21 apparatus. Flow cytometry analysis does not allow resolution of such small fragments.
22 Commercial capillary electrophoresis sequencers have sufficient resolution, but have a
23 high cost and are not suitable for rapid fragment size determination.
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1 Currently, DNA size analysis requires approximately three hours to run on a gel,
2 plus additional time to set up the sequencing unit and download the data. Flow cytometry
3 analysis does not resolve 100 to 500 base pair (bp) fragments with 1 bp resolution which
4 is required. Commercial capillary electrophoresis sequencers that can resolve 1 bp are
5 costly and do not permit size determination.

6 Capillary electrophoresis is a powerful analytical technique that can be used to
7 separate and detect a number of analytes according to their charge. Traditionally,
8 electrophoresis has been practiced on slab gels. However, the use of capillaries
9 introduces a number of advantages including small sample and electrolyte volumes, high
10 efficiency separations, and small instrument size. The capillary systems are also
11 amenable to miniaturization.

12 In capillary electrophoresis, an electrical field is applied along the length of a
13 fused silica capillary filled with a buffer. Typically, particle samples are introduced at the
14 anode and are carried towards the cathode by the electroosmotic movement of the
15 electrolyte and the electrophoretic movement of the analyte ions. The sample is separated
16 according to the charge to size ratio of the analyte ions.

17 The separated sample is detected at some optimal fixed point in the capillary. A
18 number of detection schemes are used in capillary electrophoresis including spectroscopic
19 techniques such as UV-Vis absorbance, fluorescence, and raman; mass spectrometry; and
20 electrochemical detection. Separations can be performed on analyte samples including
21 charged particles such as small organic acids and bases, proteins, peptides, amino acids,
22 and nucleic acids. Neutral molecules can also be separated with a special form of

1 capillary electrophoresis called micellar electrokinetic chromatography. Because the
2 entire sample must migrate past the detection window, the capillary electrophoretic
3 systems tend to be long and require significant time for a good separation of particles.
4

5 In light of the foregoing, it would be a significant advancement in the art to
6 provide a device that could rapidly separate and detect particles in a sample solution. It
7 would be a further advancement if the device could be produced inexpensively. It would
8 be a further advancement if the device were amenable to a high throughput system and
9 miniaturization. It would be a further advancement if the device used capillary
10 electrophoresis. Such a device is disclosed and claimed herein.
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13 14 3. BRIEF DESCRIPTION THE INVENTION

15 The present invention relates to a device and method for electrophoretic
16 separation and detection of particles. Such particles may include but are not limited to
17 nucleic acids, proteins, inorganic ions, organic ions, or neutral species. The device has a
18 separation channel in which the particles are loaded. The particles may be separated by
19 applying a charge along the length of the separation channel. After a period of time, a
20 detector may be activated to determine the location of the separated particles in the
21 separation channel.
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23

24 The separation channel may be a capillary having a first end and a second end.
25 The capillary is filled with an electrolyte buffer solution formulated for electrophoresis.
26 The buffer-filled capillary is connected adjacent the first end to a first buffer reservoir and
27 adjacent the second end to a second buffer reservoir. The buffer solution may contain
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1 tris-boric acid EDTA hereinafter, TBE, potassium tartrate, tris-acetate EDTA, hereinafter
2 TAE, and a gel sieving material. The gel may be added to mechanically separate particles
3 with a relatively constant charge to ionic radius such as oligonucleotides. Such sieving
4 materials may include but are not limited to poly(ethylene glycol), poly(vinyl alcohol),
5 hydroxy propyl methyl cellulose, hydroxyethylcellulose, linear polyacrylamide and/or
6 poly(vinylpyrrolidone). If the buffer solution has too high of a viscosity, the solution will
7 not flow properly through the capillary during loading, however, too low of a viscosity
8 may not allow optimal separation of the particles. Using the above gel sieving materials
9 in a mixed mode at a concentration in the range from about 0.1% to about 5% will create
10 a workable viscosities in the range of about 0.5 cp to about 50 cp at room temperature.

11
12 A sample of particles to be separated may be loaded adjacent the first end of the
13 capillary. An electrical source is used to apply a voltage along the capillary. Generally a
14 voltage in the range of about 4,000 V dc to about 20,000 V dc may be used. The particles
15 within the capillary migrate under the influence of the voltage in the direction of the
16 second end of the capillary. As the particles migrate they are separated.

17
18 After the voltage has been applied for a period of time, the capillary may be
19 scanned by a detector to determine the location of the separated particles. The detector
20 may scan the capillary at more than one position. Because the detector can scan more
21 than one position on the capillary, the time required to separate and detect the location of
22 the particle is reduced, compared to conventional fixed detector instruments.

23
24 The detector may use fluorescent technology to detect the particles. For
25 example, a fluorescent label may be attached to the particles prior to loading the particles
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1 in the capillary. When the particles have been separated, an excitation source such as a
2 laser, may direct an excitation beam on the capillary. The excitation source may be
3 scanned along the capillary to direct the excitation beam on more than one position along
4 the length of the capillary. The scanning of the excitation source along one or more
5 capillaries is hereinafter referred to as rastering. Rastering the beam along the part or all
6 of the length of the capillary shortens the time required to separate and detect the
7 particles. The fluorescent labels on the particle emit fluorescent light after excitation with
8 the beam.
9

11 The excitation beam may be configured to allow for optimal detection and
12 resolution of particles. An excitation beam may bleach the fluorescent tags if the beam's
13 intensity is too great. Additionally, if the beam is too weak, the fluorescent tags may not
14 emit enough light to be detected. A beam with a power in the range of about 1 mW to
15 about 1000 mW will allow the tags to fluoresce while minimizing bleaching. The
16 excitation beam may also be configured with a width sufficient to allow the light detector
17 to distinguish between particles of different sizes. A beam width in the range from about
18 5 μm to about 1000 μm can be used to excite the fluorescent tags.
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22 A light detector is positioned to detect the fluorescence and record the position of
23 the beam on the capillary when fluorescence is emitted. The position of the beam on the
24 capillary when fluorescence is detected can be used to determine the position of a
25 fluorescing particle within the capillary. The light detector may be low-level light
26 detection electronics such as photomultipliers, photodiodes, and CCD cameras. The light
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1 detector may be connected to a computer to record and determine the position of the
2 fluorescing particle.

3
4 The capillary may be covered with a coating to create a light waveguide. The
5 light waveguide retains the fluorescent light within capillary and directs the fluorescent
6 light toward the ends of the capillary. A light detector positioned adjacent either end of
7 the capillary may detect the fluorescence. A coating with low a refractive index may
8 optimally retain the fluorescent light. Generally, a coating with a refractive index
9 between about 1.1 and 1.5 is used. A coating with a refractive index of about 1.3 is
10 presently preferred. Polytetrafluoroethylene coatings such as Teflon® AF manufactured
11 by DuPont have a sufficiently low refractive index.

12
13 A fiber optic may be coupled end-on to the capillary. The fiber optic allows the
14 light detector to be positioned somewhat distantly from the end of the capillary. The
15 fiber optic can be made of a material that is optimized for transmission of the wavelength
16 of the light to be detected. Generally the wavelength of the light emitted from the
17 fluorescent tags is in the range of about 200 nm to about 1000 nm

18
19 An optical filter may be placed between the capillary and the light detection
20 electronics. The filter is selected such that the fluorescent light passing through the filter
21 is maximized while background light such as the light of the excitation laser is
22 minimized. The optical filter assembly may have a high band pass filter for filtering light
23 with a wavelength greater than about 500 nm and a notch filter for filtering the light of
24 the excitation beam. In other configurations, the optical filter may be a narrow band pass
25 filter which filters light other than the light with a wavelength corresponding to the
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1 wavelength of the light emitted from the fluorescent label ± 10 nm. Even other
2 configurations may use a light spectrometer to resolve the fluorescent light according to
3 its characteristic wavelengths and detect the resolved light.
4

5 The light detector may be configured to detect and distinguish between more
6 than one color of fluorescent light. In this manner different particles may be labeled with
7 fluorescent tags which fluoresce at different wavelengths. The detector will detect the
8 different colors of light and determine the position within the capillary of the differently
9 labeled particles. Such a configuration may be useful in such applications as sequencing
10 a nucleic acid.
11

12 The capillary typically has a length in the range from about 5 cm to about 100
13 cm. A length of less than about 20 cm may allow for sufficient separation of most
14 particle that may be used with the device. A capillary length of about 10 cm may be used
15 to create a compact capillary electrophoretic device without compromising the ability of
16 the device to separate and detect long DNA fragments.
17

18 The separation and detection device may be adapted for a high throughput
19 system by the addition of one or more capillaries. Such multi-capillary devices will allow
20 for rapid separation and identification of particles. These additional capillaries can be
21 configured and attached to the device as the single capillary discussed above. Such a
22 multi-capillary system may facilitate the sizing of particles. For example, a particle of
23 unknown size can be fluorescently labeled and loaded into a first capillary. A sample of
24 particles of known sizes can also be fluorescently labeled and loaded into a second
25 capillary. A voltage is applied along the capillaries for a period of time causing the
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1 particles to migrate within the capillaries. The capillaries are then rastered with an
2 excitation beam, and the position of the particles determined. The location of the particle
3 of unknown size may be compared to the sample of particles of a known size to
4 determine the size of the unknown. Additionally, when the device is used to sequence a
5 nucleic acid, the plurality of capillaries increases the speed at which a long nucleic acid
6 segment can be sequenced.
7

8 The present invention also relates to a method for separating and sizing of
9 particles by using the short channel capillary electrophoresis device of the present
10 invention. A sample of particles to be separated and sized is obtained and fluorescently
11 labeled. The fluorescently labeled particles are loaded into a device of the present
12 invention with one or more capillaries. A voltage is applied along one or more capillaries
13 for a period of time. The excitation beam is rastered onto the one or more capillaries.
14 The detection of the fluorescence from the excited particles is monitored and recorded.
15 The position of the excitation beam on the capillary when fluorescence is detected can be
16 used to determine the position of the particles in the capillary and relative size of the
17 particles as compared to known size standards.
18

19 The voltage is left on for a time sufficient to allow for adequate separation of the
20 particles. It has been found that a period in the range from about 8 minutes to about 18
21 minutes allows for the particles to be separated enough to be distinguished by the
22 detection system. Additionally, a period of less than about 15 minutes may be used with
23 most particles, while generally a period of about 12 minutes may be used.
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1 The device of the present invention can also be used with methods for
2 sequencing a nucleic acid. For example a sample of DNA to be sequenced is obtained.
3
4 The DNA sample is then run in standard a dideoxy sequencing reaction. The sequencing
5 reaction has four separate reaction mixtures with a different dideoxy nucleotide for each
6 standard nucleotide type. A distinct fluorescent label is added to each of the four reaction
7 mixtures. The reaction mixtures are run to form four separate reaction product. The
8 reaction products are pooled and loaded into the apparatus of the present invention with a
9 light detector capable of detecting and distinguishing between the fluorescence of the four
10 fluorescent tags. A voltage is then applied across the one or more capillaries of the
11 device. The excitation beam is rastered on the capillaries and the fluorescent light
12 collected in the light detector is monitored. A comparison of the position of the
13 excitation beam on the capillary to the color of light detected by the light detector may be
14 used to determine the position and type of a corresponding nucleotide within the DNA
15 sample, thus producing a DNA sequence.
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21 **4. DESCRIPTION OF THE DRAWINGS**

22 A more particular description of the invention briefly described above will be
23 rendered by reference to the appended drawings and graphs. These drawings and graphs
24 only provide information concerning typical embodiments of the invention and are not
25 therefore to be considered limiting of its scope.
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28 Figure 1 is a perspective view of one embodiment of a device of the present
29 invention.
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1 Figure 2 is a perspective view of an alternative embodiment of the device of the
2 present invention.

3 Figure 3 is a schematic representation of the ionic double layer formed at a
4 capillary wall constituting the major influence in the movement of bulk fluid,
5 electroosmotic flow, towards the cathode.
6

7 Figure 4 is a graph representing the separation of a 100 to 1,500 bp DNA ladder
8 using the mini-CE apparatus in a 50 cm column. Time axis in seconds.
9

10 Figure 5 is a graph representing the separation of a 1,500 bp DNA ladder
11 detected with a fixed fluorescence detector.
12

13 Figure 6 is a graph representing the detection of a 1,500 bp DNA ladder detected
14 800 seconds after start of separation by scanning the excitation laser along the length of
15 the capillary.
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18 **5. DETAILED DESCRIPTION OF THE INVENTION**
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20 The presently preferred embodiments of the present invention will be best
21 understood by reference to the drawings, wherein like parts are designated by like
22 numerals throughout. It will be readily understood that the components of the present
23 invention, as generally described and illustrated in the figures herein, could be arranged
24 and designed in a wide variety of different configurations. Thus, the following more
25 detailed description of the embodiments of the apparatus, system, and method of the
26 present invention, as represented in Figures 1 through 6, is not intended to limit the scope
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1 of the invention, as claimed, but is merely representative of presently preferred
2 embodiments of the invention

3
4 Referring to Figure 1, a capillary electrophoretic device for separating and
5 detecting particles is generally designated as 10. The device 10 can separate particles
6 including but not limited to nucleic acids, proteins, inorganic ions, organic ions, and
7 neutral species. A capillary 12 is filled with an electrolyte buffer solution 17. The
8 capillary 12 functions as a separation channel wherein the particles are electrophoretically
9 separated. A voltage is applied across the capillary 12 by an electrical source 30. As the
10 voltage is applied, the particles migrate from a first end 14 of the capillary 12 toward a
11 second end 16 of the capillary. The particles continue their migration until the electrical
12 source 30 is shut off or disconnected from the capillary 12. A detector 38 is used to scan
13 the length of the capillary 12 to detect particles within the capillary 12. The separation is
14 allowed to proceed until the first analyte reaches the prepositioned detector. The
15 separation is then stopped and the detector 38 scanned along the length of the capillary.
16 The location of the particles can be ascertained by recording the position of the detector
17 38 when the particles where detected.

18
19 The capillary 12 is dimensioned to create a compact electrophoretic device 10.
20
21 In general the capillary 12 has a length 48 in the range from about 5 cm to about 100 cm.
22
23 Capillaries 12 with a length 48 in the range from about 10 cm to about 50 cm have been
24 used to effectively separate the particles. It is presently preferred that the length 48 of the
25 capillary 12 be about 10 cm. The capillary 12 has an inside diameter in the range from
26 about 20 μm to about 150 μm . Efficient, reproducible separation and detection of the
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1 particles is possible even with these small dimensions because of the detector 38 is
2 rastered along the capillary 12.

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4 A range of voltages can be used with the electrophoretic device 10. In general
5 the voltage applied is in the range from about 4,000 V to about 20,000 V dc. A voltage
6 of about 11,000 V dc is used in one presently preferred configuration of the
7 electrophoretic device. The voltage creates a current within the capillary 12 in the range
8 from about 1 μ A to about 60 μ A. A current of about 10 μ A has been used to separate the
9 particles efficiently. Moreover, the capillary 12 does not have to be precoated to
10 deactivate the inner surface.
11

12
13 The period of time that the voltage is left on should be sufficient to allow for
14 adequate separation of the particles. It has been found that a period in the range from
15 about 8 minutes to about 18 minutes allows for the particles to be separated enough to be
16 distinguished by the detection system. Additionally, a period of less than about 15
17 minutes may be used with most particles. A period of about 12 minutes has produced
18 adequate separation of the particles.
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20

21 A first buffer reservoir 18 is in fluid communication with the first end 14 of the
22 capillary 12. A second buffer reservoir 20 is also in fluid communication with the second
23 end of the capillary 12. The buffer reservoirs 18, 20 present a convenient location for the
24 attachment of electrodes 52. As the particles migrate under the electrical current, a small
25 amount of buffer solution may also flow along the capillary. The buffer reservoirs 18, 20
26 ensure that the capillary 12 is constantly filled with buffer solution 17.
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1 The particles can be loaded into the capillary 12 by electrophoretic injection.
2 The first buffer reservoir 18 is removed and replaced with a reservoir containing a sample
3 of particles. The electrical source is activated to apply a voltage across the capillary for a
4 short period of time. A period of about 15 seconds is sufficient to cause the sample to be
5 injected into the capillary 12. Once the sample of particles is loaded, the buffer reservoir
6 18 can be replaced and the particles separated.

7
8
9 The electrolyte buffer 17 can be chosen from the various buffers generally used
10 with electrophoretic systems. The electrolyte buffer 17 is typically tris-boric acid EDTA,
11 hereinafter TBE, potassium tartrate, or tris-acetate EDTA, hereinafter, TAE. The
12 electrolyte salt is generally used at a concentration in the range of about 0.1X to about
13 2X. In a presently preferred embodiment the concentration of the electrolyte salt is about
14 0.5X.
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17 Separation of DNA fragments and other oligonucleotides presents a special
18 challenge to capillary electrophoresis because oligonucleotides have relatively constant
19 charge to ionic radius with fragment size. Thus introducing DNA fragments into free
20 zone capillary electrophoresis systems usually will result in no separation. However, the
21 use of electrophoresis as a method of transporting DNA fragments is quite advantageous
22 because flow occurs under laminar conditions with relatively constant velocity across the
23 migration zone in the capillary. Separations of DNA in a capillary electrophoretic
24 system is accomplished through mechanical sieving.
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28 In order to accomplish the mechanical separation of the DNA fragments, the
29 electrolyte buffer solution 17 further contains a gel sieving material in distilled water.
30

1 The gel sieving material is composed of cellulose polymers or their derivatives such as
2 poly(ethylene glycol), poly(vinyl alcohol), hydroxy propyl methyl cellulose (HPMC),
3 hydroxyethylcellulose, or linear polyacrylamide. The cellulose polymers or derivatives
4 are combined with a low viscosity surface coating material such as
5 poly(vinylpyrrolidone) (PVP). The sieving material should have a viscosity that allows
6 the buffer solution to be readily added and removed from the capillary while coating the
7 capillary surface. Optimally, the viscosity is in the range of about 0.5 cp to about 50 cp.
8 A viscosity of about 5 cp is used in one presently preferred embodiment. Typically the
9 sieving polymer is added at concentrations of about 0.1% to about 2%. It has been found
10 that HPMC at a concentration of about 0.5% efficiently functions as a mechanical sieve.
11 Generally, PVP is added at concentrations of about 0.1% to about 5%. A concentration
12 of about 5% PVP is presently preferred in one configuration.

13 The capillary 12 is coated with a coating 22 having a low refractive index. The
14 coating 22 transforms the capillary 12 into a light waveguide which retains light within
15 the capillary and does not exclude the excitation light from entering the capillary. The
16 light is directed toward the ends 14, 16 of the capillary. For the coating to transform the
17 capillary into a light wave guide, the coating has a refractive index number lower than the
18 refractive index of the material from which the capillary is made and lower than the
19 refractive index of the separation medium. Generally, the capillary is made from fused
20 quartz or fused silica which have a refractive index number of about 1.5. Thus, a material
21 with a refractive index number in the range of about 1.1 to about 1.4 is used. A material
22 with a refractive index number of about 1.31 is presently preferred.

1 Polytetrafluoroethylene materials such as Teflon® AF manufactured by DuPont have a
2 sufficiently low refractive index.

3
4 The detector 38 uses fluorescence detection to determine the position of the
5 particles within the capillary 12. Fluorescence detection is a sensitive detection method
6 that can be used with capillary electrophoresis. For example the separation assay
7 represented by Figure 5 was performed using TO-PRO-3 intercalating dye (642/661
8 Ex/Em). This dye can be easily excited with a small HeNe laser at 635 nm. Using TO-
9 PRO-3 allows the system to be quite sensitive, however, the sensitivity of the system can
10 be improved dramatically by using other dyes such as PicoGreen (498/520 Ex/Em).
11
12 Excitation with an argon ion laser at 488 nm may also be used, however the argon laser
13 can be bulky and cumbersome. Using diode lasers which emit at 500 nm can allow for
14 fluorescent detection of particles while using a smaller laser. Fluorescence detection is
15 generally performed by collection optics positioned at normal angles to the capillary.
16
17 High aperture optics and mirrors can be used to maximize light collection.
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19
20 As illustrated in Figure 1, an excitation source 34 such as a laser 36 or other light
21 source is positioned perpendicular to the capillary 10. The excitation source 34 is
22 mounted to a stage (not shown) which allows the excitation beam 44 to be scanned or
23 rastered in a first direction A or a second direction B along the length of the capillary 12.
24
25 The particles to be analyzed are tagged with a fluorescent label before they are injected
26 into the capillary 12. The excitation source 34 is tuned to the excitation frequency of the
27 fluorescent tag on the particles. When the excitation beam 44 contacts the fluorescent
28 labels they emit fluorescent light. Because of the coating 22, the fluorescence is
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1 contained within the capillary 12 and directed toward the ends thereof 14, 16 where the
2 light can be detected by a light detector 37. The light detector 37 is coupled to a
3 computer 54 which determines the position of a particle within the capillary 12 by
4 analyzing the position of the excitation beam 44 on the capillary 12 and the fluorescence
5 signals.
6

7 The light of the excitation beam 44 is focused on the capillary 12 to achieve the
8 maximum light intensity without causing the fluorescent tags to photo-decompose. The
9 focusing of the beam 44 on the capillary also minimizes the width of the excitation beam.
10 Because the minimized beam 44 directs light onto only a very small portion of the
11 capillary at a given time, the device 10 can detect a small separation distance thereby
12 maximizing the detection efficiency of the device 10.
13

14 The power of the excitation beam 44 can be optimized to provide maximum
15 fluorescing of the fluorescent tags while minimizing the background light transmitted
16 through the capillary 12. An excitation beam 44 with a power in the range from about 1
17 mW to about 100 mW allows for sufficient fluorescing of the tags while minimizing the
18 interfering background light within the capillary. In a presently preferred embodiment of
19 the electrophoretic device 10, the excitation beam has a power of about 30 mW.
20

21 To achieve good separation efficiencies for DNA fragments it is preferable to set
22 up a detection optics system that can allow for a separation efficiency of about 0.1 to
23 about 100 million theoretical plates. The overall achievable separation efficiency is a
24 function of the width of the excitation beam and the separation length that is being used
25 and imaged. The width of the excitation beam should be in the range from about 5 μm to
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1 about 1000 μm for a separation length of from about 5 to about 100 cm. It is presently
2 preferred that the width of the beam 44 be about 50 μm or less for a separation length of
3 about 10 cm. The excitation beam 44 is moved along the capillary 44 in a series of steps
4 small enough to distinguish between the separated particles. A step approximately equal
5 to the width of the excitation beam 44 allows for adequate distinction of particles. The
6 excitation beam 44 is moved along the capillary at a rate slow enough to maximize the
7 signal to noise ratio, yet fast enough that the separation plus detection time is minimized.
8 The detection time per laser step along the capillary is 1-1000 ms, with the preferred step
9 duration being only 5 ms.

13 A connector 24 is coupled to the capillary 12 near the second end 16 of the
14 capillary 12. The connector has a first end 23 and a second end 25. The connector 24 has
15 a channel 27 that runs from the first end 23 of the connector 24 to the second end 25.
16 The channel 27 is proportioned so that the capillary 12 fits snugly within the channel 27
17 without leaking buffer solution 17. The capillary 12 is inserted within the connector 24
18 to approximately the midpoint of the connector 24. A second channel 29 is formed in the
19 connector 24 in which a tube 39 is inserted to fluidly connect capillary 12 to the buffer
20 reservoir 20. A fiberoptic 26 is inserted within the first channel 27 to the point of almost
21 contacting the capillary 12. The fiberoptic 26 is selected to optimize the wavelength of
22 light to be detected. Generally a fiberoptic 26 optimized for light in the range from about
23 200 nm to about 1,000 nm is used. In a presently preferred configuration, the fiberoptic
24 is optimized for light with a wavelength between about 500 nm to about 600 nm.

1 A small separation distance 33 is left between the fiberoptic 26 and the capillary
2 12 to allow the capillary 12 to be in fluid communication with the buffer reservoir 20
3 while preventing the fluorescence from escaping. An optimal separation distance 33
4 depends on the numerical aperture of the capillary waveguide 12 and the fiberoptic 26.
5 Generally this separation distance 33 is in the range from about 0.1 mm to about 1.5 mm.
6 In a presently preferred embodiment, the separation distance 33 is optimized at 0.5 mm.
7 The distance 33 can be important to efficient light collection, and fluid flow of the
8 system. If the distance 33 is too large the fluorescence will not accurately be detected. If
9 the distance is too small the fluid will not flow properly.
10

11 This arrangement allows the use of a fiberoptic 26 coupled end-on to the
12 capillary 12. With this configuration, the capillary 12, with its low refractive index
13 coating, acts as a liquid core waveguide trapping and directing the fluorescent light
14 longitudinally toward the fiberoptic 26. The fiberoptic 26 directs the light into low-level
15 light detection electronics 40 which can detect and signal a computer 54 of the presence
16 of fluorescence. Suitable low-level light detection electronics 40 may include
17 photomultipliers, photodiodes or CCD cameras. Because the capillary 12 serves as a
18 light wave guide, the fiberoptic 26 and low-level light detection electronics 40 can be
19 positioned at one or both ends of the capillary 12. In the illustrated configuration, a
20 single photo multiplier 41 is used.
21

22 An optical filter 46 is positioned between the fiberoptic 26 and the light detection
23 electronics 40 to prevent incident light from the excitation source 34 from reaching the
24 detection electronics 40. The filter 46 is selected such that it maximizes the fluorescent
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1 light while minimizing the background light from the excitation beam hitting the
2 detector. The optical filter system can be a combination of a high band pass filter 46, for
3 example greater than about 500 nm, with a notch filter 46 at the wavelength of the
4 excitation beam for example about 488 nm. Alternatively, the filter 46 may be a narrow
5 band pass filter 46 at the emission wave length corresponding the fluorescence of the
6 fluorescent tags. For example a narrow band pass filter in the range of about 520 nm plus
7 or minus about 10 nm has been used. Other methods of filtering light such as a prism, or
8 light spectrometer may be also used.

11 Referring now to Figure 2, a capillary electrophoretic system for high throughput
12 analysis is generally designated 110. The high throughput is made possible by the
13 addition of one or more capillaries 112. Such multi-capillary devices 110 allow for rapid
14 separation and identification of particles. The speed of separation, detection, and sizing
15 of particles is increased because multiple samples can be run simultaneously.

18 The capillaries 112 are configured and attached to the device 110 much like the
19 single capillary 12 discussed above. The particles are loaded into multiple capillaries 112
20 and a voltage applied from an electrical source 130. After the particles are sufficiently
21 separated, the voltage is shut off and the excitation source activated to cause the particles
22 to fluoresce. The excitation source 134 can be a laser 136 or other light source as
23 discussed above. The excitation source 134 is mounted on a X-Y stage so that the single
24 beam 144 can be independently directed onto each of the capillaries 112.

28 Multiple fiberoptics 126 are coupled to the second ends 116 of the capillaries 112
29 as with the single capillary device 10. The fiberoptics 126 direct the fluorescence toward
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1 light detection electronics 140. Thus, a single photomultiplier can function to detect the
2 light from all of the capillaries 112 and direct the information into a computer 154. A
3 filter 146 can also be provided to reduce the incident light transmitted to the light
4 detection electronics 140.

6 The present invention also relates to a method for separating and sizing of
7 particles by using the short channel capillary electrophoresis device 10, 110 of the present
8 invention as illustrated in Figures 1 and 2. A sample of particles to be separated and
9 sized is obtained and fluorescently labeled. The fluorescently labeled particles are loading
10 into a device 10, 110. A voltage is applied across the one or more capillaries 12, 112 for
11 a period of time. The excitation beam 44, 144 is rastered onto the one or more capillaries
12 12, 112. The light detector 40 collects the fluorescence from the excited particles. The
13 position of the excitation beam 44, 144 on the capillary when fluorescence is detected is
14 used to determine the position of the particles in the capillary 12, 112 and the relative size
15 of the particles. A sample of particles of known size can also be loaded with the
16 unknown sample. Thus, the position of the unknown particles can be compared to the
17 position of the known particles to determine the actual size of the unknown particles.

22 The capillary electrophoretic device 10, 110 can be used to sequence DNA by
23 simply adding three more photomultipliers and filtering sets or using a light spectrometer.
24 DNA to be sequenced is run in a standard dideoxy sequencing reaction. A separate
25 reaction mixture is set up for each of the standard nucleotide types, and contains one
26 dideoxy nucleotide type. Each reaction mixture contains a distinct fluorescent label, thus
27 a label is created for each of the four standard nucleotide types. The reaction mixtures are
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1 run to form four separate reaction products. The reaction products are pooled and loaded
2 into the apparatus of the present invention 10, 110. Each of the photomultipliers is
3 selected to detect one of the four fluorescent tags. A voltage is then applied across the
4 one or more capillaries 12, 112 of the device. The excitation beam 44, 144 is rastered on
5 the capillary, and the fluorescent light is collected by the photomultipliers. A comparison
6 of the position of the excitation beam on the capillary to the color of light detected by the
7 photomultipliers is used to determine the position and type of a corresponding nucleotide
8 within the DNA sample. Photomultipliers are used because they are inexpensive and
9 very sensitive, however, other light detection systems such as CCD cameras, photo
10 diodes and light spectrometers may also be used. The fiberoptic 26 that couples the light
11 from the separation channel to the detector can be divided from the detection channel to
12 direct light to one or more light detectors.
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17 6. EXAMPLES

18 The following examples are given to illustrate various embodiments which have
19 been made with the present invention. It is to be understood that the following examples
20 are not comprehensive or exhaustive of the many types of embodiments which can be
21 prepared in accordance with the present invention.
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24 Example 1 - Electrophoresis

25 Electrophoresis takes advantage of the differences in the movement of ions in an
26 electrical field, in solution. The electrophoretic velocity (v_{ep}) for ions in a fluid with
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1 viscosity (η) and in an electrical field (E) varies according to the charge state (q) and
2 ionic radius (r) of the ion according to the equation:

$$3 \quad v_{ep} = \frac{5/3q}{\pi r \eta} E$$

6
7 Referring to Figure 3, when practiced in a fused silica capillary 210, the
8 electrophoresis process is also influenced by a bulk movement of the electrolyte fluid in
9 the capillary 210. This movement is caused by the formation of an electrical double layer
10 214 at the walls 212 of the capillary 210. The cause of this double layer 214 is based on
11 the existence of negative silanol charges 216 at the fused silica surface. These fixed
12 silanol 216 charges attract positive charges 218 from the fluid. Overall there will also be
13 an attraction of the positive charges 218 towards the cathode causing the bulk fluid
14 movement called electroosmotic (or endoosmotic) flow.
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17 The electroosmotic velocity is given by the potential arising from the charge on
18 the wall (ξ), the solution's dielectric constant (ϵ), the viscosity of the liquid (η), and the
19 electrical field (E), according to the equation:

$$20 \quad v_{eo} = \frac{\epsilon \xi}{4\pi \eta} E$$

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22 The overall velocity of the analyte in such a system is given by the summation of these
23 two vectors: $v = v_{ep} + v_{eo}$. For DNA analysis it is advantageous for the electroosmotic
24 vector to be 0 or negligible. In that way, DNA particles may migrate only under the
25 influence of their charge, which is negative, and not be opposed by the positive
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1 electroosmotic flow of the solution. This is achieved by introducing a surface
2 deactivating agent, e.g., poly(vinylpyrrolidone).
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5 Example 2 - Separation of a DNA Ladder Using TBE Buffer

6 A sample of particles was loaded into a capillary electrophoretic device of the
7 present invention. The capillary was filled with TBE buffer with a gel sieving material.
8 The gel sieving material was present at a concentration between about 0.5% and about
9 1%. The separation was performed on a device with about a 25 cm capillary length. As
10 seen in Figure 4, the separation was excellent. The separation of Figure 4 was performed
11 in about 50 minutes. However separations can be achieved in less than 10 minutes using
12 a 25 cm column but with decreased resolution at the large fragment end. Nevertheless
13 separations of PCR products that achieve +/- 5 base pair resolution at fragment lengths
14 below 500 bp, in five minutes have been achieved.
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20 Example 3 - Separation and Detection of a 1,500 bp DNA Ladder

21 A 1,500 bp DNA ladder was obtained and loaded into a standard capillary
22 electrophoretic device with a stationary detection beam. A current was applied across the
23 capillary. The particles were detected as they moved past the detection window of the
24 device. As seen in Figure 5, the separation and detection of the 1,500 bp DNA ladder
25 took about 1,400 seconds or over 23 minutes.
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28 A 1,500 bp DNA ladder was also loaded into the capillary electrophoretic device
29 of the present invention. After a period of about 13 minutes, the migration of the
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1 particles was terminated by disconnecting the voltage. The detection beam was moved
2 along the length of the capillary to image the complete electrophoretic separation of the
3 particles. In this manner, each separated fragment does not have to pass by a detection
4 window, but the complete separation channel becomes the detection window. The
5 detection process took only about 10 seconds longer of additional time. As seen in
6 Figure 6, the device of the present invention decreases the amount of time previously
7 required to separate and detect DNA fragments. The device also needed only about 13
8 cm of capillary length to fully separate the DNA ladder.
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1 **7. SUMMARY**

2 In summary, the use of multi-capillary electrophoresis technology for
3 inexpensive, sensitive and high-throughput DNA fragment and chemical sizing is
4 described. The device can separate and detect particles within a short capillary. The
5 addition of a sample of particle of known size will allow the actual size of the particles to
6 be rapidly determined. The invention is also useful for multi-color DNA sequencing
7 applications. A sample to be analyzed is introduced into one end of an electrophoresis
8 capillary, and a voltage is applied longitudinally along the capillary wall which results in
9 electrophoretic separation of the component materials in the sample. The capillary also
10 functions as a liquid-core waveguide. Detection of fluorescence resulting from the
11 interaction of light introduced through the capillary wall with the component materials is
12 then accomplished at the end of the capillary. Excitation of the separated sample
13 components in the capillary may be achieved at any point in the capillary, which permits
14 the instantaneous visualization of the entire electrophoregram. By controlling component
15 movement through the capillary, samples can be removed for further analysis using mass
16 spectrometry or DNA sequencing. The technology will allow analyses to be conducted in
17 a field laboratory without using large and expensive equipment.

18 The present invention may be embodied in other specific forms without
19 departing from its structures, methods, or other essential characteristics as broadly
20 described herein and claimed hereinafter. The described embodiments are to be
21 considered in all respects only as illustrative, and not restrictive. The scope of the
22 invention is, therefore, indicated by the appended claims, rather than by the foregoing
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description. All changes which come within the meaning and range of equivalency of the claims are to be embraced within their scope.

What is claimed and desired to be secured by United States Letters Patent is: